O'Bryen, Barbara

From:

Rabin, Evelyn

To: Subject: O'Bryen, Barbara Sequence Search 08/700737

Date:

Tuesday, June 03, 1997 8:54AM

Priority:

High

1816 305-6811

Barbara, Please search SEQ ID NOS. 12, 15, and 19. Thank you.

300 July 10

Page 1

- L7 ANSWER 5 OF 6 BIOSIS COPYRIGHT 1997 BIOSIS
- AN 94:61987 BIOSIS
- DN 97074987
- TI Differential expression in rheumatoid synovium and synovial fluid of alpha-4-beta-7 integrin. A novel receptor for Fibronectin and vascular cell adhesion molecule-1.
- AU Lazarovits A I; Karsh J
- CS Univ. Hosp., Room 4TU46, Box 5339, 339 Windermere Rd., London, ON N6A 5A5, CAN
- SO Journal of Immunology 151 (11). 1993. 6482-6489. ISSN: 0022-1767
- LA English
- AB T lymphocyte adhesion to vascular endothelium plays an important role in the immunopathogenesis of rheumatoid arthritis. The migration of T lymphocytes into the synovium is mediated by a variety of adhesion molecules, notably the integrins. We have prepared **Act**
 - 1, a murine mAb that identifies a novel integrin termed
 - alpha-4-beta-7. The natural ligands for
 - alpha-4-beta-7 are vascular cell adhesion

molecule-1 and fibronectin; both molecules are upregulated in the rheumatoid synovium. We investigated the expression of alpha -4-beta-7 in the three compartments of rheumatoid arthritis, the peripheral blood, synovial fluid, and synovial membrane, utilizing the FACS and immunoperoxidase microscopy of frozen tissues. The results of our experiments show a striking differential expression of alpha-4-beta-7 integrin in rheumatoid arthritis. Sixty-two percent of synovial membrane T cells expressed high density alpha-4-beta-7, in contrast to only 4.7% of synovial fluid and 9.1% of PBL. These data suggest that the expression of alpha-4-beta-7 integrin may provide a mechanism whereby certain T cells adhere to rheumatoid synovium while others remain in the synovial fluid. The augmented expression of alpha-4-beta-7 in the synovial membrane T cells may contribute to the development

- L5 ANSWER 1 OF 6 MEDLINE
- AN 97099142 MEDLINE
- TI Distribution of beta 7 integrins in human intestinal mucosa and organized gut-associated lymphoid tissue.
- AU Farstad I N; Halstensen T S; Lien B; Kilshaw P J; Lazarovitz A I; Brandtzaeg P
- CS Laboratory for Immunohistochemistry and Immunopathology (LIIPAT), University of Oslo, National Hospital, Norway.
- SO IMMUNOLOGY, (1996 Oct) 89 (2) 227-37. Journal code: GH7. ISSN: 0019-2805.
- CY ENGLAND: United Kingdom
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Priority Journals; Cancer Journals
- EM 9702
- EW 19970204
- Two alternative integrins involved in mucosal homing (alpha AR 4 beta 7) or epithelial retention (alpha E beta 7) of lymphocytes were examined in the human gut. The distribution of the beta 7 subunit [monoclonal antibody (mAb) M301] was bimodal in that it was strongly expressed by alpha E beta 7 + cells but weakly by alpha 4 beta 7 + cells. More than 90% of intraepithelial lymphocytes (IEL), including the minor subsets of CD4+, T-cell receptor (TCR) gamma/delta +, and CD3- cells, expressed alpha E beta 7 as did most lamina propria CD8+ (88%) and a fraction (36%) of CD4+ lymphocytes. Conversely, B-lineage cells (CD19+) and macrophages (CD68+) were negative. In gut-associated lymphoid tissue (GALT: Peyer's patches and appendix) only a few (< 5%) cells were positive for alpha E beta 7 (confined to CD8+ lymphocytes and CD11c+ putative dendritic cells). A relatively small fraction of IEL (30-50%) expressed alpha 4 beta 7 (mAb Act-1), while most (70%) lamina propria T and B lymphocytes, blasts, plasma cells and macrophages were positive. In GALT, T lymphocytes expressed similar levels of alpha 4 beta 7 as in the lamina propria whereas relatively few B lymphocytes (< 50%) were positive. Isolated lamina propria CD8+, CD4+, CD19+, and CD38+ cells contained mRNA for alpha 4 and the former three subsets as well as appendix CD8+ cells also for beta 7 while only lamina propria CD8+ cells had mRNA for alpha E. Together, the results suggested that alpha E beta 7 and alpha 4 beta 7 are differentially regulated in inductive sites and effector sites of the human gut. Because lymphoid cells at both sites expressed mainly alpha 4 beta 7, this integrin may be a homing receptor on memory and effector cells bound for lamina propria as well as on naive lymphocytes extravasating in GALT. Conversely, because alpha E beta 7 was mainly expressed by CD8+ cells in epithelium and lamina propria, it was probably induced after extravasation, in agreement with the observation that IEL and a fraction of lamina propria T lymphocytes (mainly CD8+ cells) generally expressed higher levels of beta 7 than most CD4+ and B cells. Also a subset of putative dendritic cells located near the follicle-associated epithelium of GALT expressed alpha E beta 7, perhaps reflecting epithelial

interaction during primary immune responses.

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ANSWER 2 OF 6 MEDLINE
L5
AN
     97067800
                  MEDLINE
     Integrin alpha 4 beta 7 mediates human
ΤI
     eosinophil interaction with MAdCAM-1, VCAM-1 and fibronectin.
     Walsh G M; Symon F A; Lazarovils A L; Wardlaw A J
ΑU
     Department of Respiratory Medicine University of Leicester Medical
     School, Glenfield General Hospital, UK.
     IMMUNOLOGY, (1996 Sep) 89 (1) 112-9.
so
     Journal code: GH7. ISSN: 0019-2805.
     ENGLAND: United Kingdom
CY
     Journal; Article; (JOURNAL ARTICLE)
DT
LA
     English
FS
     Priority Journals; Cancer Journals
EM
     9702
EW
     19970204
AΒ
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We have investigated the contribution of integrin alpha 4 beta 7 to human peripheral blood eosinophil adhesive interactions. Immunofluorescence and flow cytometry demonstrated constitutive expression of alpha 4 beta 7 by eosinophils. Expression of alpha 4 beta 7 or alpha 4 beta 7 was not enhanced by eosinophil activation with platelet-activating factor (PAF). Expression of alpha 4 beta 7 was confirmed by

immuno-precipitation of 125I-labeled lysates analysed by sodium dodecyl sulphate polyacrylamide gel electrophoresis (SDS PAGE). Approximately 20% of unstimulated eosinophils were adherent to L1-2 cells transfected with vascular cell adhesion molecule-1 (VCAM-1) cDNA, while very few resting eosinophils adhered to mouse mucosal adressin cell adhesion molecule-1 (MAdCAM-1) transfectants. Binding of unstimulated eosinophils to VCAM-1 transfectant was inhibited by HPI 2 (an antibody that blocks both alpha 4 beta

1 and alpha 4 beta 7 functions), but not

Act-1, and alpha 4 beta 1

monoclonal antibody (mAb). PAF stimulation resulted in increased binding of eosinophils to MAdCAM-1 transfectants, which was inhibited by both HPI 2 and Act-1. In contrast, PAF did not enhance binding to VCAM 1 transfectants, although binding of PAE-stimulated eosinophils to VCAM-1 could be partially inhibited by Act-1. Stimulation of eosinophils with the beta 7-activating mAb TS2 16 resulted in enhanced binding of eosinophils to both VCAM-1 and MAdCAM-1 transfectants. The increased binding was largely alpha 4 beta 7-dependent. Unstimulated eosinophils bound to soluble recombinant human (rh) VCAM-1 and fibronectin (Fn), coated on 96-well plates in dose-dependent manner. Binding was inhibited by HPI-2 and 4b4, an anti-beta 1 mAb, but not by Act-1. TS2 16 treatment increased adherent cell numbers and this enhanced binding was inhibited by Act-1. We have therefore confirmed that alpha 4 beta 7 is functionally active on unstimulated eosinophils. In contrast, PAF-induced enhancement of eosinophils binding to VCAM-1 or MAdCAM-1 was alpha 4 beta 7-dependent. In addition treatment with TS2 16 resulted in a alpha 4 beta 7-dependent enhancement of eosinophil binding to VCAM-1, MAdCAM-1 and Fn. We therefore hypothesize that alpha 4 beta 7 may have an important role in eosinophil localization in

diseases such as asthma and inflammatory bowel disease.

- L5 ANSWER 3 OF 6 MEDLINE
- AN 97067798 MEDLINE
- TI Expression and function of alpha 4/beta 7 integrin on human natural killer cells.
- AU Perez-Villar J J; Zapata J M; Melero I; Postigo A; Sanchez-Madrid E; Lopez-Botet M
- CS Servn to de Immunologia Hospital de la Prineesa, Madrid, Spain.
- SO IMMUNOLOGY, (1996 Sep) 89 (1) 96-104. Journal code: GH7. ISSN: 0019-2805.
- CY ENGLAND: United Kingdom
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Priority Journals; Cancer Journals
- EM 9702
- EW 19970204
- AB In this report we have analysed the expression and function of the alpha 4/beta 7 heterodimer in human natural killer (NK) cells. The expression of alpha 4 beta 7 is induced in NK cells upon activation as the anti alpha 4 beta 2 ACT-1 monoclonal antibody (mAb) family stained a minority of peripheral blood NK cells, whereas it

family stained a minority of peripheral blood NK cells, whereas it strongly reacted with in vitro long-term interleukin-2

(IL-2)-activated NK cells. Incubation with ACT-1

on its F(ab) fragments induced a strong homotypic adhesion of NK cells, comparable to than stimulated by the anti-alpha

4 HPI 7 mAb. Cell cell interaction induced by the

ACT-1 mAb was only prevented by another anti-

alpha 4 mAb (HP2.1) that recognizes a different

epitope. In alpha 4 beta 7-mediated cell

aggregation the alpha 4 beta 7 heterodimer was

redistributed to intercellular contact sites thus, suggesting a direct involvement of this integrin in the formation of cell clusters. In NK cells attached to Fibronectin (FN38) or vascular cell adhesion molecule-1 (VCAM-1), both alpha 4

beta 7 and alpha 4 beta 7 integrins were

redistributed at the ventral cellular membrane forming discrete contact sites. The ACT-1 mAb only partially

blocked NK cell binding to FN38, but in combination with the anti-beta 7 mAb LIAI 2, NK cell binding to FN38 was completely inhibited. In contrast. **ACT-1** did not modify NK

cell adhesion to VCAM-1 thus supporting the theory that the alpha 4 beta 7 binding sites for both ligands

appear to be different. Our results indicate that upon

IL-2-activation, expression of functional alpha 4

beta-integrin is induced on NK cells potentially participating in their interaction with both extracellular matrix and endothelial cells.

- L5 ANSWER 4 OF 6 MEDLINE
- AN 96286047 MEDLINE
- TI Specific inhibition of T lymphocyte coactivation by triggering integrin beta 1 reveals convergence of beta 1, beta 2, and beta 7 signaling pathways.
- AU Woodside D G; Teague T K; McIntyre B W
- CS Department of Immunology, University of Texas, M.D. Anderson Cancer Center, Houston 77030, USA.
- NC CA62596 (NCI)
- SO JOURNAL OF IMMUNOLOGY, (1996 Jul 15) 157 (2) 700-6.

Journal code: IFB. ISSN: 0022-1767.

- CY United States
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Abridged Index Medicus Journals; Priority Journals; Cancer Journals
- EM 9701
- EW 19970104
- AB T cell coactivation is a dynamic process subject to integrin-dependent positive and negative regulation. Costimulation of human peripheral blood T cells by CD3 mAb OKT3 in conjunction with anti-alpha 4 has been shown to be down-regulated by the anti-beta 1.1 epitope-specific mAb 18D3. As expected, maximal costimulation induced by alpha 4 -specific mAb L25 was inhibited (70%) by the addition of soluble mAb 18D3. Surprisingly, soluble mAb 18D3 inhibited maximal proliferation induced by the costimulatory alpha 4 beta 7-specific mAb ACT-1 by 40%, thus demonstrating that one integrin subfamily can regulate the activity of another. To determine whether mAb 18D3 could regulate more than alpha
 - 4-associated integrin-mediated costimulation, nonalpha 4 integrins were tested. mAb 18D3 inhibited maximal proliferation induced by alpha 4
 - -specific mAb 3D6, and an alpha 4-specific mAb
 - 16. This clearly demonstrates that a variety of integrin costimulatory molecules (of the beta 1, beta 2, and beta 7 subfamilies) can be regulated negatively by mAb 18D3. To analyze the specificity of this negative regulation, other cell surface costimulatory molecules were tested for susceptibility to mAb 18D3. Although Abs specific for CD4, CD26, CD28, CD44, CD45RA, or CD45RO were sufficient to activate T cells when co-immobilized with anti-CD3 mAb, all were refractory to the inhibitory effects of mAb 18D3. Inhibition of T cell activation directly correlated with diminished IL-2 production. This suggests that mAb 18D3 selectively regulates integrin-dependent T cell activation by delivering a negative effect at some common point utilized by various integrin subfamilies.
- L5 ANSWER 5 OF 6 MEDLINE
- AN 95261715 MEDLINE
- TI Integrin alpha 4 beta 7 co-stimulation of human peripheral blood T cell proliferation.
- AU Teaque T K; Lazarovits A I; McIntyre B W
- CS Department of Immunology, University of Texas M. D. Anderson Cancer, Center, Houston 77030, USA.
- NC CA62596 (NCI)
- SO CELL ADHESION AND COMMUNICATION, (1994 Dec) 2 (6) 539-47.

 Journal code: B4A. ISSN: 1061-5385.
- CY Switzerland
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Priority Journals
- EM 9508
- AB The integrin alpha 4 beta 7 mediates lymphocytes adhesion to VCAM-1 on activated endothelium, fibronectin in the extracellular matrix, and the mucosal vascular addressin MAdCAM-1. It is unclear whether alpha 4 beta 7 performs any function beyond directing specific adhesion reactions. We addressed the possibility that triggering of alpha 4 beta 7 with a specific monoclonal antibody was capable of

delivering an accessory stimulus that would coactivate T cells and lead to proliferation. At submitogenic levels of anti-CD3 stimulation, triggering of alpha 4 beta 7 by immobilized mAb ACT-1 resulted in T cell blastogenesis, IL-2 production, expression of the IL-2 receptor alpha chain CD25, and ultimately DNA synthesis. These results indicate that the integrin alpha 4 beta 7 is involved in more than lymphocyte adhesion and homing but also plays a role in cell signaling.

- L5 ANSWER 6 OF 6 MEDLINE
- AN 93329067 MEDLINE
- TI Selective expression of integrin alpha 4 beta 7 on a subset of human CD4+ memory T cells with Hallmarks of gut-trophism.
- AU Schweighoffer T; Tanaka Y; Tidswell M; Erle D J; Horgan K J; Luce G E; Lazarovits A I; Buck D; Shaw S
- CS Experimental Immunology Branch, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892.
- NC AR60684 (NIAMS) HL07185 (NHLBI)
- SO JOURNAL OF IMMUNOLOGY, (1993 Jul 15) 151 (2) 717-29. Journal code: IFB. ISSN: 0022-1767.
- CY United States
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Abridged Index Medicus Journals; Priority Journals; Cancer Journals
- EM 9310

. . . .

AB Human memory CD4+ T lymphocytes are heterogenous in expression of integrins; one subset has the unexpected phenotype beta 1 low alpha 4 high. We demonstrate that this subset is unique among CD4+ cells in expression of high levels of alpha 4 beta 7, detected by a distinctive mAb Act-1. alpha 4 beta 7 is

involved in binding to both fibronectin and vascular cell adhesion molecule-1; Act-1 blocks cell binding to the former and augments binding to the latter. Act-1 expression marks a subset of memory cells that, unlike the predominant circulating memory cell, has up-regulated beta 7 rather than beta 1. Their phenotype is distinct from that described for skin-homing T cells and is fully consistent with that described for gut-homing T cells. Differential adhesion capacity of this subset is verified by selective binding to FN and vascular cell adhesion molecule-1 in a beta 1-independent fashion. Thus, alpha 4 beta 7 detected on this subset of circulating normal T cells fits the expectations for a gut-homing receptor.

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Logon file001 12jul99 17:37:04
ANNOUNCEMENT
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NEW
***Market Guide Company Financials (File 100)
***Frost & Sullivan Market Engineering (File 767)
***Canada Newswire (File 616 for current news, File 816 for archive)
***So America Bus Info (File 617 for current news, File 817
     for archive news)
***UPI News (Files 261 for current news & 861 for archive news)
***Africa News (Files 606 for current news & 806 for archive news)
***ITAR/TASS (Files 607 for current news & 667 for archive news)
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           of new databases, price changes, etc.
***** The DIALORDER suppliers DYNAMIC and FILEDOC are no longer
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***** details, please contact MARUZEN CO. LTD, at 3-3272-3496.
***** Jupiter Communications removed May 14.
***** Preliminary records through 05/05
*** File 332 is currently unavailable. ***
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*File 399: Use is subject to the terms of your user/customer agreement.
RANK charge added; see HELP RATES 399.
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*File 357: Derwent changes DialUnit pricing from May 1, 1999. See
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292 BETA7

33 ALPHA4(W)BETA7

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S2 3 RD S1 (unique items)

? t s2/7/all

2/7/1 (Item 1 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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09144254 97376878

Structure-function analysis of the integrin beta 7 subunit: identification of domains involved in adhesion to MAdCAM-1.

Tidswell M; Pachynski R; Wu SW; Qiu SQ; Dunham E; Cochran N; Briskin MJ; Kilshaw PJ; Lazarovits AI; Andrew DP; Butcher EC; Yednock TA; Erle DJ

Lung Biology Center, Department of Medicine, University of California, San Francisco 94143, USA. easd@itsa.ucsf.edu

J Immunol (UNITED STATES) Aug 1 1997, 159 (3) p1497-505, ISSN 0022-1767 Journal Code: IFB

Contract/Grant No.: K08HL03230, HL, NHLBI; R01HL52004, HL, NHLBI; AI37832, AI, NIAID

Languages: ENGLISH

Document type: JOURNAL ARTICLE

Beta 7 integrins serve special roles in mucosal immunity. Alpha 4 beta 7-mediated adhesion to mucosal addressin cell adhesion molecule-1 (MAdCAM-1) directs lymphocyte homing to the gut, and alpha E beta 7 mediates binding of lymphocytes to E-cadherin on epithelial cells. Since alpha 4 beta 7 mediates adhesion to MAdCAM-1 but alpha 4 beta 1 does not, we used beta 7/beta 1 chimeras to directly assess the importance of specific regions of beta 7 in MAdCAM-1 binding. We found a region of beta 7 (residues 46-386) that accounts for specificity of alpha 4 beta 7 binding to MAdCAM-1. We also used human/mouse and human/rat chimeric beta $\tilde{7}$ subunits to map epitopes recognized by fifteen anti-beta 7 mAbs. Six of seven Abs that block adhesion to MAdCAM-1 and E-cadherin (Fib 21, 22, 27, 30, 504; Act-1) mapped to amino acid residues 176-250. Residues 176-250 lie within the region of beta 7 that specifies MAdCAM-1 binding and also within a region that has a predicted structure homologous to the metal ion-dependent adhesion site (MIDAS) domains of the integrin subunits alpha L and alpha M. Three new Abs that recognize beta 7 in the presence of Mn2+, but not Ca2+, and promote adhesion to MAdCAM-1, mapped to amino acids 46-149. One blocking and five other Abs mapped to other regions (amino acids 387-725). We conclude that a MIDAS-like domain serves a critical role in beta 7 integrin-mediated adhesion.

2/7/2 (Item 2 from file: 155) DIALOG(R)File 155:MEDLINE(R)

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08839966 97099142

Distribution of beta 7 integrins in human intestinal mucosa and organized gut-associated lymphoid tissue [published erratum appears in Immunology 1997 Jun; 91(2):322]

Farstad IN; Halstensen TS; Lien B; Kilshaw PJ; Lazarovits AI; Brandtzaeg P; Lazarovitz AI [corrected to Lazarovits AI]

Laboratory for Immunohistochemistry and Immunopathology (LIIPAT), University of Oslo, National Hospital, Norway.

Immunology (ENGLAND) Oct 1996, 89 (2) p227-37, ISSN 0019-2805

Journal Code: GH7

Languages: ENGLISH

Document type: JOURNAL ARTICLE

Two alternative integrins involved in muco'sal homing (alpha 4 beta 7) or epithelial retention (alpha E beta 7) of lymphocytes were examined in the human gut. The distribution of the beta 7 subunit [monoclonal antibody (mAb) M301] was bimodal in that it was strongly expressed by alpha E beta 7 + cells but weakly by alpha 4 beta 7 + cells. More than 90% of intraepithelial lymphocytes (IEL), including the minor subsets of CD4+, T-cell receptor (TCR) gamma/delta +, and CD3- cells, expressed alpha E beta 7 as did most lamina propria CD8+ (88%) and a fraction (36%) of CD4+ lymphocytes. Conversely, B-lineage cells (CD19+) and macrophages (CD68+) were negative. In gut-associated lymphoid tissue (GALT: Peyer's patches and appendix) only a few (< 5%) cells were positive for alpha E beta 7 (confined to CD8+ lymphocytes and CD11c+ putative dendritic cells). A relatively small fraction of IEL (30-50%) expressed alpha 4 beta 7 (mAb Act-1), while most (70%) lamina propria T and B lymphocytes, blasts, plasma cells and macrophages were positive. In GALT, T lymphocytes expressed similar levels of alpha 4 beta 7 as in the lamina propria whereas relatively few B lymphocytes (< 50%) were positive. Isolated lamina propria CD8+, CD4+, CD19+, and CD38+ cells contained mRNA for alpha 4 and the former three subsets as well as appendix CD8+ cells also for beta 7 while only lamina propria CD8+ cells had mRNA for alpha E. Together, the results suggested that alpha E beta 7 and alpha 4 beta 7 are differentially regulated in inductive sites and effector sites of the human gut. Because lymphoid cells at both sites expressed mainly alpha 4 beta 7, this integrin may be a homing receptor on memory and effector cells bound for lamina propria as well as on naive lymphocytes extravasating in GALT. Conversely, because alpha E beta 7 was mainly expressed by CD8+ cells in epithelium and lamina propria, it was probably induced after extravasation, in agreement with the observation that IEL and a fraction of lamina propria T lymphocytes (mainly CD8+ cells) generally expressed higher levels of beta 7 than most CD4+ and B cells. Also a subset of putative dendritic cells located near the follicle-associated epithelium of GALT expressed alpha E beta 7, perhaps reflecting epithelial interaction during primary immune responses.

2/7/3 (Item 3 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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08806160 96286047

Specific inhibition of T lymphocyte coactivation by triggering integrin beta 1 reveals convergence of beta 1, beta 2, and beta 7 signaling pathways.

Woodside DG; Teague TK; McIntyre BW

Department of Immunology, University of Texas, M.D. Anderson Cancer Center, Houston 77030, USA.

J Immunol (UNITED STATES) Jul 15 1996, 157 (2) p700-6, ISSN 0022-1767 Journal Code: IFB

Contract/Grant No.: CA62596, CA, NCI

Languages: ENGLISH

Document type: JOURNAL ARTICLE

T cell coactivation is a dynamic process subject to integrin-dependent positive and negative regulation. Costimulation of human peripheral blood T cells by CD3 mAb OKT3 in conjunction with anti-alpha 4 has been shown to be down-regulated by the anti-beta 1.1 epitope-specific mAb 18D3. As expected, maximal costimulation induced by alpha 4-specific mAb L25 was inhibited (70%) by the addition of soluble mAb 18D3. Surprisingly, soluble mAb 18D3 inhibited maximal proliferation induced by the costimulatory alpha 4 beta 7-specific mAb ACT-1 by 40%, thus demonstrating that one integrin subfamily can regulate the activity of another. To determine whether mAb 18D3 could regulate more than alpha 4-associated integrin-mediated costimulation, non-alpha 4 integrins were tested. mAb 18D3 inhibited maximal proliferation induced by alpha 4-specific mAb 3D6, and an alpha 4-specific mAb 16. This clearly demonstrates that a variety of integrin costimulatory molecules (of the beta 1, beta 2, and beta 7

subfamilies) can be regulated negatively by mAb 18D3. To analyze the specificity of this negative regulation, other cell surface costimulatory molecules were tested for susceptibility to mAb 18D3. Although Abs specific for CD4, CD26, CD28, CD44, CD45RA, or CD45RO were sufficient to activate T cells when co-immobilized with anti-CD3 mAb, all were refractory to the inhibitory effects of mAb 18D3. Inhibition of T cell activation directly correlated with diminished IL-2 production. This suggests that mAb 18D3 selectively regulates integrin-dependent T cell activation by delivering a negative effect at some common point utilized by various integrin subfamilies.

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            1875 ALPHA4
             292 BETA7
            1875 ALPHA4
             292 BETA7
              33 ALPHA4 (W) BETA7
             637 ANTIBOD? AND (ALPHA4 OR BETA7 OR ALPHA4(W)BETA7)
? s s3 and (alpha4(w)beta7 or alpha4beta7)
             637
                  s3
            1875 ALPHA4
             292 BETA7
              33 ALPHA4 (W) BETA7
             204 ALPHA4BETA7
      S4
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11967567
           BIOSIS NO.: 199900220880
Jejuna of patients with insulin-dependent diabetes mellitus (IDDM) show
  signs of immune activation.
AUTHOR: Savilahti E(a); Ormala T; Saukkonen T; Sandini-Pohjavuori U;
  Kantele J M; Arito A; Ilonen J; Akerblom H K
AUTHOR ADDRESS: (a) Hospital for Children and Adolescents, University of
  Helsinki, Stenbackinkatu 11, FIN-00290, Hel, Finland
JOURNAL: Clinical and Experimental Immunology 116 (1):p70-77 April, 1999
ISSN: 0009-9104
DOCUMENT TYPE: Article
RECORD TYPE: Abstract
LANGUAGE: English
SUMMARY LANGUAGE: English
           (Item 2 from file: 5)
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11828952
          BIOSIS NO.: 199900075061
The development of experimental autoimmune encephalomyelitis in the mouse
  requires alpha4-integrin but not alpha4beta7-integrin.
```

AUTHOR: Engelhardt Britta(a); Laschinger Melanie; Schulz Martina; Samulowitz Ulrike; Vestweber Dietmar; Hoch Gabi AUTHOR ADDRESS: (a)Max-Planck Institut Physiologische, Klinische Forschung, W.G. Kerck-Hoff-Institut, Abt. Molekula, Germany

JOURNAL: Journal of Clinical Investigation 102 (12):p2096-2106 Dec. 15,

1998

ISSN: 0021-9738

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

5/3/3 (Item 3 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
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11782775 BIOSIS NO.: 199900028884

MAdCAM-1 costimulates T cell proliferation exclusively through integrin alpha4beta7, whereas VCAM-1 and CS-1 peptide use alpha4beta1:
Evidence for "remote" costimulation and induction of hyperresponsiveness to B7 molecules.

AUTHOR: Lehnert Klaus; Print Cristin G; Yang Yi; Krissansen Geoffrey W(a) AUTHOR ADDRESS: (a) Univ. Auckland, Sch. Med., Park Rd., Grafton, Auckland, New Zealand

JOURNAL: European Journal of Immunology 28 (11):p3605-3615 Nov., 1998

ISSN: 0014-2980

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

5/3/4 (Item 4 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 1999 BIOSIS. All rts. reserv.

11610774 BIOSIS NO.: 199800392539 Blockade of both L-selectin and **alpha4** integrins abrogates naive CD4 cell trafficking and responses in gut-associated lymphoid organs.

AUTHOR: Bradley Linda M(a); Malmo Mary E; Fong Sherman; Tonklonogy Susan L; Watson Susan R

AUTHOR ADDRESS: (a) Dep. Immunol., Scripps Res. Inst., IMM-23, 10550 North Torrey Pines, Road, La Jolla, CA 92037, USA

JOURNAL: International Immunology 10 (7):p961-968 July, 1998

ISSN: 0953-8178

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

5/3/5 (Item 1 from file: 73)
DIALOG(R)File 73:EMBASE
(c) 1999 Elsevier Science B.V. All rts. reserv.

07694830 EMBASE No: 1999172498

Lymphocyte migration in lymphocyte function-associated antigen (LFA)-1-deficient mice

Berlin-Rufenach C.; Otto F.; Mathies M.; Westermann J.; Owen M.J.; Hamann A.; Hogg N.

N. Hogg, Leukocyte Adhesion Laboratory, Imperial Cancer Research Fund,

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Lincoln's Inn Fields, London WC2A 3PX United Kingdom
 AUTHOR EMAIL: hogg@icrf.icnet.uk
 Journal of Experimental Medicine ( J. EXP. MED. ) (United States) 03 MAY
 1999, 189/9 (1467-1478)
                ISSN: 0022-1007
  CODEN: JEMEA
  DOCUMENT TYPE: Journal; Article
  LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH
 NUMBER OF REFERENCES: 51
           (Item 2 from file: 73)
 5/3/6
DIALOG(R) File 73: EMBASE
(c) 1999 Elsevier Science B.V. All rts. reserv.
            EMBASE No: 1999059648
07579020
  Cooperative activity of alpha4beta1 and alpha4beta7 integrins in
mediating human B-cell lymphoma adhesion and chemotaxis on fibronectin
through recognition of multiple synergizing binding sites within the
central cell-binding domain
  Yin Z.; Giacomello E.; Gabriele E.; Zardi L.; Aota S.-I.; Yamada K.M.;
Skerlavaji B.; Doliana R.; Colombatti A.; Perris R.
  Dr. R. Perris, Division for Experimental Oncology 2, Ctro. di Riferimento
 Oncol. Aviano, Istituto Nazionale Centroeuropeo, Aviano (PN) 33081 Italy
 AUTHOR EMAIL: rperris@ets.it
 Blood ( BLOOD ) (United States) 15 FEB 1999, 93/4 (1221-1230)
                ISSN: 0006-4971
  CODEN: BLOOA
 DOCUMENT TYPE: Journal; Article
  LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH
 NUMBER OF REFERENCES: 33
           (Item 3 from file: 73)
 5/3/7
DIALOG(R)File 73:EMBASE
(c) 1999 Elsevier Science B.V. All rts. reserv.
07088726
             EMBASE No: 1997370590
 Mucosal immunity in the female genital tract
  Brandtzaeg P.
  P. Brandtzaeg, Lab. Immunohistochemistry (LIIPAT), Institute of
  Pathology, University of Oslo, N-0027 Oslo 1 Norway
 AUTHOR EMAIL: per.brandtzag@rh.uio.no
  Journal of Reproductive Immunology ( J. REPROD. IMMUNOL. ) (Ireland)
  1997, 36/1-2 (23-50)
                ISSN: 0165-0378
  CODEN: JRIMD
  PUBLISHER ITEM IDENTIFIER: S0165037897000612
  DOCUMENT TYPE: Journal; Review
  LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH
 NUMBER OF REFERENCES: 108
           (Item 4 from file: 73)
 5/3/8
DIALOG(R) File 73: EMBASE
(c) 1999 Elsevier Science B.V. All rts. reserv.
             EMBASE No: 1997298776
07011841
  Fibronectin type III5 repeat contains a novel cell adhesion sequence,
KLDAPT, which binds activated alpha4beta1 and alpha4beta7 integrins
  Moyano J.V.; Carnemolla B.; Dominguez-Jimenez C.; Garcia-Gila M.; Albar
J.P.; Sanchez-Aparicio P.; Leprini A.; Querze G.; Zardi L.; Garcia-Pardo A.
  A. Garcia-Pardo, CIB, CSIC, Velazquez 144, 28006 Madrid Spain
  AUTHOR EMAIL: cibgp96@fresno.csic.es
  Journal of Biological Chemistry ( J. BIOL. CHEM. ) (United States) 1997
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CODEN: ANYAA ISSN: 0077-8923 DOCUMENT TYPE: Journal; Conference Paper LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

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(Item 8 from file: 73)
DIALOG(R) File 73: EMBASE
(c) 1999 Elsevier Science B.V. All rts. reserv.
            EMBASE No: 1996313472
 Phenotype, and migration properties of three major subsets of tissue
homing T cells in sheep
 Mackay C.R.; Andrew D.P.; Briskin M.; Ringler D.J.; Butcher E.C.
  LeukoSite Inc., 215 First Street, Cambridge, MA 02142 United States
  European Journal of Immunology (EUR. J. IMMUNOL. ) (Germany) 1996,
  26/10 (2433-2439)
               ISSN: 0014-2980
  CODEN: EJIMA
  DOCUMENT TYPE: Journal; Article
  LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH
           (Item 9 from file: 73)
 5/3/13
DIALOG(R) File 73: EMBASE
(c) 1999 Elsevier Science B.V. All rts. reserv.
            EMBASE No: 1996249706
06585083
  The role of alpha4 integrins in lung pathophysiology
  Lobb R.R.; Pepinsky B.; Leone D.R.; Abraham W.M.
  Biogen Inc, 14 Cambridge Center, Cambridge, MA 02142 United States
  European Respiratory Journal, Supplement (EUR. RESPIR. J. SUPPL. ) (
  Denmark) 1996, 9/22 (104S-108S)
               ISSN: 0904-1850
  CODEN: ERJSE
  DOCUMENT TYPE: Journal; Conference Paper
                     SUMMARY LANGUAGE: ENGLISH
  LANGUAGE: ENGLISH
            (Item 10 from file: 73)
 5/3/14
DIALOG(R) File 73: EMBASE
(c) 1999 Elsevier Science B.V. All rts. reserv.
             EMBASE No: 1996026641
06377271
  In vivo migration of radiolabelled lymphocytes in rheumatoid synovial
tissue engrafted in SCID mice: Implication of beta2 and beta7
-integrin
  Jorgensen C.; Couret I.; Hellier I.; Bologna C.; Canovas F.; Brochier J.;
Reme T.; Sany J.
  Service d'Immuno-Rhumatologie, Centre Gui-de-Chauliac, 34295 Montpellier
  Cedex 5 France
  Journal of Rheumatology ( J. RHEUMATOL. ) (Canada) 1996, 23/1 (32-35)
  CODEN: JRHUA ISSN: 0315-162X
  DOCUMENT TYPE: Journal; Article
  LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH
            (Item 11 from file: 73)
 5/3/15
DIALOG(R) File 73: EMBASE
(c) 1999 Elsevier Science B.V. All rts. reserv.
             EMBASE No: 1995240062
  Distinc roles of L-selectin and integrins alpha4beta7 and LFA-1 in
lymphocyte homing to Peyer's patch-HEV in situ: The multistep model
confirmed and refined
  Bargatze R.F.; Jutila M.A.; Butcher E.C.
  Lab. of Immunology/Vascular Biology, Department of Pathology, Stanford
  Univ. School of Medicine, Stanford, CA 94305 United States
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Immunity ( IMMUNITY ) (United States) 1995, 3/1 (99-108)
                ISSN: 1074-7613
 CODEN: IUNIE
 DOCUMENT TYPE: Journal; Article
                    SUMMARY LANGUAGE: ENGLISH
 LANGUAGE: ENGLISH
            (Item 12 from file: 73)
 5/3/16
DIALOG(R) File 73: EMBASE
(c) 1999 Elsevier Science B.V. All rts. reserv.
            EMBASE No: 1995237575
06210427
  Adhesion molecule expression and adhesion properties of murine intestinal
intraepithelial lymphocyte hybridomas
 Ni J.; Hollander D.; Sydora B.; Panwala C.
  Human Genome Sciences, Inc., 9620 Medical Center Drive, Rockville, MD
  20850-3338 United States
  Cellular Immunology ( CELL. IMMUNOL. ) (United States) 1995, 164/1
  (156-160)
                 ISSN: 0008-8749
  CODEN: CLIMB
  DOCUMENT TYPE: Journal; Article
  LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH
            (Item 13 from file: 73)
 5/3/17
DIALOG(R) File 73: EMBASE
(c) 1999 Elsevier Science B.V. All rts. reserv.
             EMBASE No: 1995227069
06203831
  Construction and adhesive properties of a soluble MAdCAM-1-Fc chimera
expressed in a baculovirus system: Phylogenetic conservation of
receptor-ligand interaction
  Yang Y.; Sammar M.; Harrison J.E.B.; Lehnert K.; Print C.G.; Leung E.;
Prestidge R.; Krissansen G.W.
  Department of Molecular Medicine, School of Medicine, University of
  Auckland, Auckland New Zealand
  Scandinavian Journal of Immunology ( SCAND. J. IMMUNOL. ) (United Kingdom
  ) 1995, 42/2 (235-247)
                 ISSN: 0300-9475
  CODEN: SJIMA
  DOCUMENT TYPE: Journal; Article
                      SUMMARY LANGUAGE: ENGLISH
  LANGUAGE: ENGLISH
            (Item 14 from file: 73)
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DIALOG(R) File 73: EMBASE
(c) 1999 Elsevier Science B.V. All rts. reserv.
             EMBASE No: 1995230437
06199138
  The alpha4 integrin chain is a ligand for alpha4beta7 and
alpha4beta1
  Altevogt P.; Hubbe M.; Ruppert M.; Lohr J.; Von Hoegen P.; Sammar M.;
Andrew D.P.; McEvoy L.; Humphries M.J.; Butchers E.C.
  Tumor Immunology Programme, German Cancer Research Center, Im Neuenheimer
  Feld 280, D-69120 Heidelberg Germany
  Journal of Experimental Medicine ( J. EXP. MED. ) (United States) 1995,
  182/2 (345-355)
  CODEN: JEMEA
                 ISSN: 0022-1007
  DOCUMENT TYPE: Journal; Article
  LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH
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5/3/19 (Item 15 from file: 73)

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DIALOG(R) File 73: EMBASE
 (c) 1999 Elsevier Science B.V. All rts. reserv.
              EMBASE No: 1995109880
 06079396
   Lymphocytes infiltrating the CNS during inflammation display a
 distinctive phenotype and bind to VCAM-1 but not to MAdCAM-1
   Engelhardt B.; Conley F.K.; Kilshaw P.J.; Butcher E.C.
   Department of Pathology, Lab Immunology and Vascular Biology, Stanford
   Univ School of Medicine, Stanford, CA 94305 United States
   International Immunology (INT. IMMUNOL.) (United Kingdom) 1995, 7/3
   (481 - 491)
   CODEN: INIME
                  ISSN: 0953-8178
   DOCUMENT TYPE: Journal; Article
   LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH
             (Item 16 from file: 73)
  5/3/20
 DIALOG(R) File 73: EMBASE
 (c) 1999 Elsevier Science B.V. All rts. reserv.
              EMBASE No: 1995052113
   Identification of putative ligand-binding sites of the integrin
 alpha4beta1 (VLA-4, CD49d/CD29)
   Kamata T.; Puzon W.; Takada Y.
   Department of Vascular Biology, The Scripps Research Institute, 10666
   North Torrey Pines Road, La Jolla, CA 92037 United States
   Biochemical Journal (BIOCHEM. J.) (United Kingdom) 1995, 305/3
   (945 - 951)
   CODEN: BIJOA ISSN: 0264-6021
   DOCUMENT TYPE: Journal; Article
   LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH
             (Item 17 from file: 73)
  5/3/21
 DIALOG(R) File 73: EMBASE
 (c) 1999 Elsevier Science B.V. All rts. reserv.
              EMBASE No: 1995009367
   Dual binding capacity of mucosal immunoblasts to mucosal and synovial
 endothelium in humans: Dissection of the molecular mechanisms
   Salmi M.; Andrew D.P.; Butcher E.C.; Jalkanen S.
   National Public Health Institute, Kiinamyllynkatu 13,20520 Turku Finland
   Journal of Experimental Medicine ( J. EXP. MED. ) (United States) 1995,
   181/1 (137-149)
   CODEN: JEMEA ISSN: 0022-1007
   DOCUMENT TYPE: Journal; Article
   LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH
  5/3/22
             (Item 18 from file: 73)
 DIALOG(R) File 73: EMBASE
. (c) 1999 Elsevier Science B.V. All rts. reserv.
              EMBASE No: 1994342969
  The pathophysiologic role of alpha4 integrins in vivo
  Lobb R.R.; Hemler M.E.
  Biogen, Inc., 14 Cambridge Center, Cambridge, MA 02142 United States
  Journal of Clinical Investigation ( J. CLIN. INVEST. ) (United States)
  1994, 94/5 (1722-1728)
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CODEN: JCINA ISSN: 0021-9738 DOCUMENT TYPE: Journal; Review

DIALOG(R) File 73: EMBASE (c) 1999 Elsevier Science B.V. All rts. reserv. EMBASE No: 1994201028 05792337 Identification of a combinatorial epitope expressed by the integrin alpha4betal heterodimer involved in the regulation of cell adhesion Bednarczyk J.L.; Szabo M.C.; Wygant J.N.; Lazarovits A.I.; McIntyre B.W. Dept. of Immunology, Texas Univ. M.D. Anderson Can. Ctr., Box 180, 1515 Holcombe Blvd., Houston, TX 77030 United States Journal of Biological Chemistry (J. BIOL. CHEM.) (United States) 1994 , 269/11 (8348-8354) ISSN: 0021-9258 CODEN: JBCHA DOCUMENT TYPE: Journal; Article LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH 5/3/24 (Item 20 from file: 73) DIALOG(R) File 73: EMBASE (c) 1999 Elsevier Science B.V. All rts. reserv. EMBASE No: 1993264514 05496415 alpha4beta7 Integrin mediates B cell binding to fibronectin and vascular cell adhesion molecule-1: Expression and function of alpha4 integrins on human B lymphocytes Postigo A.A.; Sanchez-Mateos P.; Lazarovits A.I.; Sanchez-Madrid F.; De Landazuri M.O. Servicio de Inmunologia, Hospital de la Princesa, C/Diego de Leon 62,28006 Madrid Spain Journal of Immunology (J. IMMUNOL.) (United States) 1993, 151/5 (2471 - 2483)CODEN: JOIMA ISSN: 0022-1767 DOCUMENT TYPE: Journal; Article LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH 5/3/25 (Item 21 from file: 73) DIALOG(R) File 73:EMBASE (c) 1999 Elsevier Science B.V. All rts. reserv. 05451804 EMBASE No: 1993219903 alpha4beta7 Integrin mediates lymphocyte binding to the mucosal vascular addressin MAdCAM-1 Berlin C.; Berg E.L.; Briskin M.J.; Andrew D.P.; Kilshaw P.J.; Holzmann B.; Weissman I.L.; Hamann A.; Butcher E.C. Lab. of Immunology/Vascular Biology, Department of Pathology, Stanford University, Stanford, CA 94305 United States Cell (CELL) (United States) 1993, 74/1 (185-195) CODEN: CELLB ISSN: 0092-8674 DOCUMENT TYPE: Journal; Article LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH 5/3/26 (Item 22 from file: 73)

DIALOG(R) File 73: EMBASE (c) 1999 Elsevier Science B.V. All rts. reserv.

EMBASE No: 1992155449 05015233 Role of integrin alpha4beta7/alpha4betaP in lymphocyte adherence to fibronectin and VCAM-1 and in homotypic cell clustering
Ruegg C.; Postigo A.A.; Sikorski E.E.; Butcher E.C.; Pytela R.; Erle D.J.
Department of Medicine, Lung Biology Center, University of California, San
Francisco, CA 94143 United States
Journal of Cell Biology (J. CELL BIOL.) (United States) 1992, 117/1
(179-189)

CODEN: JCLBA ISSN: 0021-9525 DOCUMENT TYPE: Journal; Article

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

5/3/27 (Item 1 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 1999 Dialog Corporation. All rts. reserv.

09963407 99265740

Integrins alpha4beta7 and alphaEbeta7 are expressed on epidermotropic T cells in cutaneous T cell lymphoma and spongiotic dermatitis.

Schechner JS; Edelson RL; McNiff JM; Heald PW; Pober JS

Department of Dermatology, Yale University School of Medicine, New Haven, Connecticut 06520-8059, USA.

Lab Invest (UNITED STATES) May 1999, 79 (5) p601-7, ISSN 0023-6837 Journal Code: KZ4

Contract/Grant No:: T32-AR07016-23, AR, NIAMS; R37-HL36003, HL, NHLBI

Languages: ENGLISH

Document type: JOURNAL ARTICLE

5/3/28 (Item 2 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 1999 Dialog Corporation. All rts. reserv.

09775096 99043885

Differences in immune responses induced by oral and rectal immunizations with Salmonella typhi Ty21a: evidence for compartmentalization within the common mucosal immune system in humans.

Kantele A; Hakkinen M; Moldoveanu Z; Lu A; Savilahti E; Alvarez RD; Michalek S; Mestecky J

University of Alabama at Birmingham, Birmingham, Alabama, USA. anu.kantele@ksshp.fi

Infect Immun (UNITED STATES) Dec 1998, 66 (12) p5630-5, ISSN 0019-9567 Journal Code: GO7

Contract/Grant No.: AI 28147, AI, NIAID; DE 01882, DE, NIDR; AI 34970, AI, NIAID; +

Languages: ENGLISH

Document type: JOURNAL ARTICLE

5/3/29 (Item 3 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 1999 Dialog Corporation. All rts. reserv.

08994387 97193612

Adhesion of multiple myeloma peripheral blood B cells to bone marrow fibroblasts: a requirement for CD44 and alpha4beta7.

Masellis-Smith A; Belch AR; Mant MJ; Pilarski LM

Department of Oncology, University of Alberta, Edmonton, Canada.

Cancer Res (UNITED STATES) Mar 1 1997, 57 (5) p930-6, ISSN 0008-5472

Journal Code: CNF Languages: ENGLISH

Document type: JOURNAL ARTICLE

5/3/30 (Item 4 from file: 155)

DIALOG(R)File 155:MEDLINE(R)

(c) format only 1999 Dialog Corporation. All rts. reserv.

08947560 97167595

Bone marrow fibroblast exposure to the inflammatory cytokines tumor necrosis factor-alpha and interferon-gamma increases adhesion of acute myeloid leukemia cells and alters the adhesive mechanism.

Bendall LJ; Kortlepel K; Gottlieb DJ

Department of Haematology, University of Sydney, Australia.

Exp Hematol (UNITED STATES) Feb 1997, 25 (2) p132-9, ISSN 0301-472X

Journal Code: EPR Languages: ENGLISH

Document type: JOURNAL ARTICLE

5/3/31 (Item 5 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 1999 Dialog Corporation. All rts. reserv.

08930257 97146023

Homing potentials of circulating lymphocytes in humans depend on the site of activation: oral, but not parenteral, typhoid vaccination induces circulating **antibody** -secreting cells that all bear homing receptors directing them to the gut.

Kantele A; Kantele JM; Savilahti E; Westerholm M; Arvilommi H; Lazarovits A; Butcher EC; Makela PH

National Public Health Institute, Helsinki, Finland.

J Immunol (UNITED STATES) Jan 15 1997, 158 (2) p574-9, ISSN 0022-1767 Journal Code: IFB

Contract/Grant No.: AI37832, AI, NIAID

Languages: ENGLISH

Document type: JOURNAL ARTICLE

5/3/32 (Item 6 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 1999 Dialog Corporation. All rts. reserv.

08689961 96202509

Stimulation of tyrosine phosphorylation after ligation of **beta7** and beta1 integrins on human B cells.

Manie SN; Astier A; Wang D; Phifer JS; Chen J; Lazarovits AI; Morimoto C; Freedman AS

Department of Medicine, Harvard Medical School, Boston, MA, USA.
Blood (UNITED STATES) Mar 1 1996, 87 (5) p1855-61, ISSN 0006-4971

Journal Code: A8G

Contract/Grant No.: CA55207, CA, NCI

Languages: ENGLISH

Document type: JOURNAL ARTICLE